A peptide comprising or consisting of the sequence YMVH or MVHW or VHWK and having at least 70% homology with part or all of the sequence EPHRWSSYMVHWK. A mixture of the peptide of claim 12 or claim 13 with another peptide having at least 4 amino acid residues and having at least 70% homology with the eta-amyloid precursor sequence DAEFRHDSGYEVHHOK. /3
A probe consisting of the peptide of claim 1/2 or claim , labelled with a signal moiety, or immobilised on a support A probe consisting of the peptide of claim 1/4, labelled with a signal moiety, or immobilised on a support. A compound which competes with the peptide of claim 12 or claim is for binding to a receptor therefor and which thereby inhibits the biological activity of the said peptide. A compound as claimed in claim 1 , wherein the biological activity is modulating a calcium-charmel-opening activity. A compound as claimed in claim if, which is capable of crossing the blood-brain barrier. An antibody to the peptide of claim 1/2 or claim 13. An antibody as claimed in claim 20 which is of the IgG class. An antibody fragment or chimeric or humanised antibody

comprising variable regions of the antibody of claim 20.

A method of treating a patient suffering from a disorde of the central nervous system or stroke or cancer, which method comprises administering to the patient a compound according to claim 14. A method of treating a patient suffering from a disorder of the central nervous/system or stroke or cancer, which method comprises administering to the patient an antibody according to claim 20. method of controlling cytoplasmic calcium concentration in vive, which method comprises administering a compound according to claim 1%. method \of controlling / cytoplasmic concentration in vivo, which method comprises administering an antibody according to claim 20. A peptide as claimed in claim 12 or claim 13, which pertide contains no more than about 14 amino acid residues, A peptide as claimed in claim 1/2 or claim 1/8, which ı, -peptide does not form part of a larger protein having homology with ij

the AChE molecule.

29. A peptide as claimed in claim 18 or claim 18 which peptide is a fragment of the AChE molecule.

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A peptide as claimed in claim 12 or claim 1/3, which peptide has been chemically synthesised. --